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(54) Title: GENE EXPRESSION CASSETTE CONTAINING NON-CODING SEQUENCE OF GROWTH HORMONE GENE

#### (57) Abstract

The present invention provides a genetic expression cassette for use in obtaining expression of a cDNA sequence in animal cells. The expression cassette comprises an inducible promoter and the 3' non-coding sequence of exon 5 of the growth hormone gene or a portion thereof. The cDNA sequence is inserted between the inducible promoter and the exon 5 of the growth hormone genes.

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#### GENE EXPRESSION CASSETTE CONTAINING NON-CODING SEQUENCE OF GROWTH HORMONE GENE

#### FIELD OF THE INVENTION

The present invention relates to a gene expression cassette which enables expression of cDNA sequences in 5 animal cells. The expression cassette of the present invention is particularly useful in achieving high-level expression of bacterial and/or plant genes in animal cells. BACKGROUND OF THE INVENTION

It is now possible to transfer unique pieces of DNA 10 between organisms in such a way that the transferred material becomes a functional part of the genetic information of the recipient organisms. The animals that are produced by this technique are termed "transgenic". One application of this technology is to transfer 15 biochemical pathways from bacteria to domestic animals in order to increase animal productivity. One difficulty which is frequently encountered in efforts to produce such transgenic animals is the lack, or very low levels of expression of the transferred DNA sequences.

The present inventors have developed a genetic expression cassette which provides information for the expression of heterologous genes, in particular bacterial genes, in mammalian cells and in several tissues of transgenic animals, at levels that provide ready detection 25 of the encoded polypeptides.

The expression cassette consists of two components:a regulatory element and a non-coding sequence from the growth hormone gene.

#### SUMMARY OF THE PRESENT INVENTION

30 Accordingly, in a first aspect the present invention consists in a genetic expression cassette for use in obtaining expression of a cDNA sequence in animal cells, the cassette comprising an inducible promoter and the 3' non-coding sequence of exon 5 of the growth hormone gene or a portion thereof, the cDNA sequence being positioned

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between the inducible promoter and the 3' non-coding sequ no of exon 5 of the growth hormone gene.

In a preferred embodiment of the present invention the inducible promoter is the immediate upstream nucleotide sequence of the sheep metallothionein-Ia gene.

The expression cassette of the present invention provides a means for the expression of a wide range of genes in transgenic animals, including the coding sequences of bacterial enzymes, plant chitinases, insecticidal scorpion venom toxin and the insecticidal protein of the bacteria <u>Bacillus thuringiensis</u>. In a preferred embodiment of the present invention the cDNA sequence is selected from the group consisting of <u>cysE</u>, <u>cysK</u>, <u>aceA</u> and <u>aceB</u> genes of <u>Escherichia coli</u> and the coding sequences of plant chitinases.

In yet a further preferred embodiment of the present invention the genetic expression cassette has a sequence substantially as shown in Figure 1.

The expression cassette of the present invention is useful in obtaining high levels of expression of cDNA sequences in animal cells. Accordingly, in a second aspect the present invention consists in a non-human animal including the genetic expression cassette of the first aspect of the present invention.

In a preferred embodiment of this aspect the animal is ovine or bovine.

# DETAILED DESCRIPTION OF THE INVENTION

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In order that the nature of the present invention may be more clearly understood, preferred forms thereof will now be described with reference to the following examples and figures in which:-

Figure 1 shows the nucleotide sequence of the expression cassette of the present invention;

Figure 2 shows the sequence of MTCE10;

35 Figure 3 shows the sequence of MTCK7;

Figure 4 shows the sequence of MTCEK1;

Figure 5 shows the sequence of MTAceA2;

Figure 6 shows the sequence of MTAceB2;

Figure 7 shows the sequence of MTAceAB11; and

Figure 8 shows levels of radiolabelled cysteine in transgenic mice containing MTCEK1 (——————————) and in control mice (- - - -). The arrow shows the position of cysteic acid.

Initially, a number of gene arrangements for 10 expression of the <u>cysK</u> gene in murine L-cells were trialled. The trialled constructs were as follows:-

pMTCK7 - sheep metallothionein-Ia gene promoter - <a href="mailto:cysK">cysK</a> - exon 5 of sheep growth hormone.

pMTCK8 - sheep metallothionein-Ia promoter - exon 1

15 sheep growth hormone - cysk - exon 5 sheep growth hormone.

pMTCK11 - sheep metallothionein-Ia promoter - <a href="mailto:cysk">cysk</a> - whole sheep growth hormone.

pMTCK12 - sheep metallothionein-Ia - exon 1 sheep growth hormone - cysk - exons 2, 3, 4 and 5 sheep growth hormone.

The constructs were transfected into murine L-cells and the O-acetylserine sulfhydrylase activity of the transfected cells measured. The results obtained are set out in Table 1.

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#### TABLE 1

# O-Acetylserine Sulfhydrylase Activity in Transfected Murine L-Cells Using Various cysk Genes

	<u>Gene</u>	Enzyme Activity
30		(nMoles cysteine produced/mg protein/30 min)
	pMTCK7	1350 <u>+</u> 24
	pMTCK8	510 <u>+</u> 13
	pMTCK11	162 <u>+</u> 17
	pMTCK12	159 <u>+</u> 6

35 (values represent the means of two determinations)

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As can seen from these results exon 5 of the growth hormone gene of sheep is required for optimum expression of genes inserted into the cassette. Other combinations which comprise larger portions of the sheep growth hormone gene are less effective in providing expression.

Two examples of the function of the expression cassette are shown as follows:

 Expression of the cysE and cysK genes of E. coli in transgenic animals

In order to provide a pathway for the biosynthesis of the amino acid cysteine, the coding sequences for the bacterial enzymes serine transacetylase and 0-acetylserine sulfhydrylase have been inserted into the expression cassette.

Three genes are described. Genes 1 and 2 each encode single bacterial proteins, gene 1 encoding the protein serine transacetylase and gene 2 encoding the protein O-acetylserine sulfhydrylase. Gene 3 is a compound gene constructed from gene 1 and gene 2, and encodes both the serine transacetylase protein and the O-acetylserine sulfhydrylase protein.

The expression cassette of the present invention was produced using methods well known in the art. Briefly this involves the steps of:

- 25 1. Isolation and cloning of the sheep metallothionein-Ia promoter sequence.
  - 2. Isolation and modification of the bacterial coding sequence and fusion to the bacterial coding sequence.
- 3. Fusion of exon 5 of the sheep growth hormone gene to 30 the metallothionein promoter/bacterial coding sequence complex.

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In order to provide further details on construction of the cassette the procedure followed in construction of MTCE10 was as follows: Step 1.

A bacterial plasmid containing the sheep metallothionein-Ia gene was digested with the restriction enzymes Eco RI and BamHl and a DNA fragment encoding the promoter region of the gene separated by agarose gel electrophoresis and cloned in the plasmid vector pUC8.

10 Step 2.

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The coding sequence and associated 5' and 3' DNA encompassing the cysE gene of Escherichia coil was cloned in the plasmid vector pGEM3 as an Eco R1 fragment excised from a lambda transducing phage containing portion of the Sub-fragments of this insert were then 15 <u>E.coil</u> chromosome. cloned into the bacteriophage M13 and the clones encompassing the bacterial initiation codon and the bacterial stop codon were used for site-directed mutagenesis to introduce a Bam H1 site at the 5' end of 20 the coding sequence and a Sau 3A site at the 3' end of the The mutagenesis was carried out on single-strand DNA by conventional procedures and the resulting modified DNA used to replace the corresponding DNA fragments in the insert of the original pGEM3 clone. A Bam H1 - Sau 3A 25 fragment of DNA was then excised from this plasmid and inserted into a similarly digested sample of the plasmid containing the metallothionein-Ia sequence. Step 3.

The plasmid containing the metallothionein-Ia

30 promoter-csyE coding sequence was digested with Pvu II (adjacent to the introduced Sau 3A site) and to this was ligated a blunt-ended Pst 1 DNA fragment isolated from the sheep growth hormone gene and encompassing exon 5.

Plasmids containing the correct orientation of the growth hormone sequence were identified by restriction enzyme mapping.

#### GENE DETAILS

Gene 1 (MTCE10)

This gene consists of the sheep metallothionein-Ia gene promoter sequence joined to the coding sequence of 5 the Escherichia coli cysE gene at a unique BamH1 restriction enzyme site. This sequence was then joined to the 3' sequence of exon 5 of the sheep growth hormone gene. Minor sequence modification in the vicinity of the initiation and stop codons of the bacterial cysE gene were 10 made by site-directed mutagenesis using synthetic oligonucleotides. The metallothionein promoter replaces all regulatory sequences located 5' to the cysE gene coding sequence, and the growth hormone exon 5 sequence replaces all untranslated sequences located 3' to the cysE gene coding sequence. The gene is approximately 3580 base 15 pairs in length, of which 2827 nucleotides have been The sequence of gene 1 is shown in Figure 2. sequenced.

Gene 2 (MTCK7)

This gene consists of the sheep metallothionein-Ia 20 gene promoter sequence joined to the coding sequence of the Escherichia coli cysK gene at a unique Sal 1 restriction enzyme site. This sequence was then joined to the 3' sequence of exon 5 of the sheep growth hormone gene. Minor sequence modification of the cysk gene in the vicinity of the initiation codon was made by site-directed mutagenesis using a synthetic oligonucleotide. metallothionein promoter replaces all regulatory sequences located 5' to the cysk coding sequence, and the sheep growth hormone exon 5 replaces all untranslated sequence located 3' to the cvsK coding sequence. The size of the gene is approximately 3750 base pairs in length, of which 2957 base pairs have been sequenced. The sequence of gene 2 is shown in Figure 3.

Gene 3 (MTCEK1)

35 This gene consists of a fusion of genes 1 and 2 to

create a single DNA sequence that encodes both the serin transacetylase and the O-acetylserine sulfhydrylase enzymes. Each coding sequence is separately regulated by its own adjacent sheep metallothionein-Ia gene promoter sequence, and each coding sequence is separately followed by the 3' sequence of exon 5 of the sheep growth hormone gene. The gene is approximately 7550 base pairs in size, of which 5784 nucleotides have been sequenced. The sequence of gene 3 is shown in Figure 4.

# 10 Example 2. The expression of the glyoxylate cycle in transgenic animals

In order to provide the enzymes needed for the operation of the glyoxylate cycle in transgenic animals, the <u>E. coli</u> genes encoding the enzymes isocitrate lyase and malate synthase have been inserted into the expression cassette.

Three genes are described. Genes 1 and 2 each encode single bacterial proteins, gene 1 encoding the protein isocitrate lyase and gene 2 encoding the protein malate synthase. Gene 3 is a compound gene constructed from gene 1 and gene 2, and encodes both the isocitrate lyase and the malate synthase proteins.

#### GENE DETAILS

#### Gene 4 (MTAceA2)

This gene consists of the sheep metallothionein-Ia gene promoter sequence joined to the coding sequence of the Escherichia coli aceA gene at a unique BamHl restriction enzyme site. This sequence was then joined to the 3' sequence of exon 5 of the sheep growth hormone

30 gene. Minor sequence modification in the vicinity of the initiation and stop codons of the bacterial aceA gene were made by site-directed mutagenesis using synthetic oligonucleotides. The metallothionein promoter replaces all regulatory sequences located 5' to the aceA gene

35 coding sequence, and the growth hormone exon 5 sequence

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replaces all untranslated sequences located 3' to the <u>aceA</u> gene coding sequence. The gene is approximately 3580 base pairs in length, of which 2827 nucleotides have been sequenced. The sequence of gene 4 is shown in Figure 5.

Gene 5 (MTAceB2)

This gene consists of the sheep metallothionein-Ia gene promoter sequence joined to the coding sequence of the Escherichia coli aceB gene at a unique Sal 1 restriction enzyme site. This sequence was then joined to the 3' sequence of exon 5 of the sheep growth hormone gene. Minor sequence modification of the aceB gene in the vicinity of the initiation codon was made by site-directed mutagenesis using a synthetic oligonucleotide. The metallothionein promoter replaces all regulatory sequences located 5' to the aceB coding sequence, and the sheep growth hormone exon 5 sequence replaces all untranslated sequence located 3' to the aceB coding sequence. The size of the gene is approximately 3750 base pairs in length, of which 2957 base pairs have been sequenced. The sequence

# Gene 6 (MTAceAB1)

This gene consists of a fusion of genes 1 and 2 to create a single DNA sequence that encodes both the isocitrate lyase and the malate synthase enzymes. Each coding sequence is separately regulated by its own adjacent sheep metallothionein-Ia gene promoter sequence, and each coding sequence is separately followed by the 3' sequence of exon 5 of the sheep growth hormone gene. The gene is approximately 7550 base pairs in size, of which 5784 nucleotides have been sequenced. The sequence of gene 6 is shown in Figure 7.

# REGULATION OF THE GENES

Regulation in Cultured Cells

Genes 1 to 6 have been transfected into mouse L-cells

in cultur to produc stably transformed cell lines. The xpression of ach gene was measured by:

- Northern blot analysis of extracted RNA.
- 2. Enzyme assay of cell extracts.

An RNA transcript of the expected size was detected in RNA extracted from each cell line, using a probe specific for the appropriate coding sequence of each gene. The intensity of the hybridisation increased when cells were grown in a medium containing 10 uM zinc sulphate, indicating that the genes were regulated by heavy metals.

The results of enzyme assays of cell extracts from each of the transformed cell lines are shown in Table 1 (genes 1 - 3) and Table 4 (genes 4,5). High levels of activity of serine transacetylase, 0-acetylserine sulfhydrylase, isocitrate lyase and malate synthase were measured in the appropriate cell extracts, and the enzyme levels were increased when cells were grown in zinc-supplemented growth media.

20 Cell extracts prepared from cells containing the fusion gene MTCEK1 contained both serine transacetylase and O-acetylserine sulfhydrylase enzyme activities, indicating that both coding sequences within the fusion gene were transcribed and translated. Furthermore, when extracts from this cell line were incubated with the substrates serine and H<sub>2</sub>S, substantial quantities of cysteine were produced, evidence that the entire biochemical pathway is operational in these cells. Similarly, cell extracts prepared from the cells containing the fusion gene MTACeAB1 contained both isocitrate lyase and malate synthase enzyme activities, indicating that both coding sequences within the fusion gene were transcribed and translated.

Expression in Transgenic Mice

35 Genes 1 to 6 were each transferred to transgenic mice

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by the technique of single-cell embryo pronuclear microinjection. Mice containing the new genes were analyzed for expression by extracting mRNA and preparing cell-free supernatants from various tissues including liver, kidney and intestine. As shown in Tables 3 and 5, high levels of activity of the various enzymes were detected in appropriate transgenic mice. Furthermore, the expression of the genes in the intestinal tissues was highly zinc-dependent.

# 10 <u>TABLE 2</u> Expression of MTCE10 and MTCK7 in transformed mouse L-cells <u>Serine Transacetylase O-acetylserine</u>

				<u>Sulfhydrylase</u>	
	cells	-Zn.	+Zn	-Zn	+zn
15	control	0	0	0	0
	MTCE10	1281	2706	_	-
	MTCK7	_		38	1367
	MTCEK1	120	360	1082	7790
				£	:- /20 -:

20 Values are nmoles product formed/mg protein/30 min

TABLE 3
Activity of serine transacetylase (SAT) and O-acetylserine sulphydrylase (OAS) in tissue extracts prepared from transgenic mice. CK7-26 contains the gene pMTCK7, CE10-29 contains pMTCE10 and CEK1-28 and CEK1-8 contains pMTCEK1. Specific activity is measured as nmoles substrate utilised (SAT) or product formed (OAS/30 min/mg protein.

	MOUSE LINE	ORGAN	SAT	<u>OAS</u>
	CK7-26	Intestine		206
10		Kidney	-	352
	•	Liver	***	13
	CE10-29	Intestine	6,546	<del>-</del> .
		Kidney	0	
		Liver	0	-
15	CEK1-28	Intestine	1,161	2,797
		Kidney	0	24
		Liver	0	3
		Brain	16	86
	CEK1-8	Intestine	4,522	12,778
20		Kidney	105	128
		Liver	9	3
		Brain	· <b>O</b>	245
			0	158
		Skin	0	329
25			6	295

In order to assess the ability of transgenic mic containing the pMTCEK1 gene to produce cysteine, transgenic mice including this gene and control mice were given 25 mm  $znso_d$  in their drinking water for a minimum 5 of four days. On the day of the experiment the  $ZnSO_4$ was relaced with normal drinking water and 60 min. later 30 - 60 uCi of Na $_2^{35}$ S was administered per os. mice were sacrificed 60 min. later and intestinal tissue homogenised in a buffered aqueous solution containing 10mM 10 dithiothreitol. Two volumes of performic acid were then added and the solution left at room temperature overnight. The suspension was then extracted with chloroform/methanol by conventional means and the aqueous layer concentrated by evaporation. Aliquots of the 15 solution were then placed on Whatman 3mm filter paper and subjected to electrophoresis in a solution of pyridine:acetic acid:H20 (10:100:900, pH3.6) at a voltage of 200 Volts for 2 hr. The paper was the cut into 0.5 cm strips and radioactivity counted in a scintillation 20 counter under standard conditions. The results are shown in Figure 8. As can be seen from these results the transgenic mice were able to synthesise radiolabelled cysteine from the administered sodium sulphide in contrast to the control mice.

#### 25 TABLE 4

Expression of MTAceA2 and MTAceB2 in transformed mouse L-cells

	cell line	isocitrate	lyase	malate	synthase
	control	0		0	
30	MTAceA2	68		-	
	MTAceB2	-		3	4.3

Values are nmoles product/mg protein/20 min

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TABLE 5
Expression of MTAceAB1 in transgenic mice

Mouse	<u>Tissue</u>	<u>Isocitrate Lyase</u>	Malate Synthase
control	intestine	not detectable	not detectable
	liver	not detectable	not detectable
	kidney	not detectable	not detectable
MTAceAB1	intestine	27.2	ND
	liver	not detectable	182
	kidney	not detectable	1.6

Values of isocitrate lyase are nmoles product/mg protein/20 min, and for malate synthase are picomoles product/mg protein/20 min (x  $10^{-2}$ )

It will be appreciated by persons skilled in the art that numerous variations and/or modifications may be made to the invention as shown in the specific embodiments without departing from the spirit or scope of the invention as broadly described. The present embodiments are, therefore, to be considered in all respects as illustrative and not restrictive.

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#### CLAIMS: -

- A genetic expression cassette for use in obtaining expression of a cDNA sequence in animal cells, the cassette comprising an inducible promoter and the 3' 5 non-coding sequence of exon 5 of the growth hormone gene or a portion thereof, the cDNA sequence being positioned between the inducible promoter and the 3' non-coding sequence of exon 5 of the growth hormone gene.
- A genetic expression cassette as claimed in claim 1 10 in which the inducible promoter is the immediate upstream nucleotide sequence of the sheep metallothionein-Ia gene.
  - A genetic expression cassette as claimed in claim 1 or claim 2 in which the cDNA codes for a bacterial enzyme, plant chitinase, insecticidal scorpion vermon toxin or the insecticidal protein of Bacillus thuringiensis.
  - A genetic expression cassette as claimed in claim 3 in which the cDNA sequence is selected from the group consisting of cysE, cysK, aceA and aceB genes of Escherichia coli.
- A genetic expression cassette as claimed in claim 1 20 5. in which the expression cassette has a sequence substantially as shown in Figure 1.
  - A transgenic non-human animal including the genetic expression cassette as claimed in any one of claims 1 to 5.
- A transgenic non-human animal as claimed in claim 6 25 in which the animal is ovine or bovine.

#### FIG. $1 \frac{1}{2}$

#### SEQUENCE OF THE EXPRESSION CASSETTE

metallothionein promoter gaattcaaagaggaaaagtgatgaaacaaggcttggcacagactccctggtatgtaattc 61 tcaggactattcaaagggaaatacccactgtcttacttcgttattggatgccagctctgc 121 ccatcacttacaaggatgcttttcctagggggcatcctatgactagggaacctccatcct ggagccgggtggactggctaggcagtggattccctggcccattcatctattcagtcgtgg agaatgtaaggaaggctgggcgacagaaggctgagttcgctgctggggctgttacaggaga aactagagactctgttcaaagtccagggtgggggctgtggggggaaatattagggaagcg gggttcgggggataggtggtgaagctcacatccatcacgggtctctgcacacgacacagg 421 qqctccaqccaaqcctqggatqtqaqcacqaggctcggattqcqcatqaqctctqggaaa 481 gggtgaaagcaaagactgcgggggggaaggactgcgaggactcagggactgg 601 teetggagegetaettgteattegggaeaaagteeeteegegttgggggggagtaggggg 661 acggaggcgtttcggtgcgcacggagcccagccgcgttccgggaatcttgcgctcggccg cgcgtggtgctcaccgcccgacccgggtgcagcgggcagctcgggtgcaggcgggggcag 781 metallothionein cap site accetetgegeeeggeeegeeteetgtgggtataatagegeteggeteetgggeteeaae 841 acgcctcccaccggaccagtggatccaca INSERT GENE IN THIS POSITION 910 growth hormone exon 5 tgtcctgtgatctaatgtcctgtgatcccgctgcgccttctagttgcca 960 gccatctgctgttacccctccctgtgccttcctagaccctggaaggtgccactccagtgc 1020 ccaccgtcctttcttaataaagcggaggaaattgcatcacattgtctgagtaggtgtcat 1080 1140 aggggtgctgtgggctctatgggtacccaggtgctgaataattgacccggttcctcctgg 1200 ggcagaaagaagcaggcacatccccttctctgtgacacacccggtcctcgcccctggtcc 1260 ttagttccagccccactcataggacactcacagctcaggagggctccgccttcaatccca cccgctaaagtgcttggagcggtctctccctctcagccaccagccgaatctaggcctcca

2040

# 2/25 FIG. 1 2/2

1380 gagtgggaagaatttaagcaagacaggctatgaagtacagagggagagaaaatgcctcca acatgtgaggaagtgatgagagaaagcgtagaattagttttgtggcataaattttaaggt 1560 tecagetetttgtgaceceaeggactgtggctgceaggetectctgtccatgggattete 1620 cagggcaagaatactggaggggttgccattccccaggggatcttcccagcccaaggatc 1680 aaacccgagtttctgcattgcaggcagattctttactctctgagccatcagggaagccct 1740 gtgggaaatgggaaccatgcaagaatggctttgggaccaataggaccagaatgtttggga 1800 tctgaactgggtcaagagatgtggaagagattctaaatgcatgtgttcatgctaagtg 1860 gcttcagtcgtgtcctactatttgcaaccccgatgaactgcagccaccaggctcctctgt 1920 1980 aacccaqqqattqaccaqqatctcttqtatctcctqgcacttqacaqqcaaatctctcac

cactagcgccactggacccagtctaag--unsequenced region

#### FIG. 2 1/3

SEQUENCE OF THE MTCE10 GENE metallothionein promoter gaattcaaagaggaaaagtgatgaaacaaggcttggcacagactccctggtatgtaattc 61 tcaqqactattcaaaqqqaaatacccactqtcttacttcqttattqqatqccaqctctqc 121 ccatcacttacaaqqatqcttttcctaggqqqcatcctatqactaqqqaacctccatcct 181 ggagccgggtggactggctaggcagtggattccctggcccattcatctattcagtcgtgg 241 agaatgtaaggaaggctgggcgacagaaggctgagttcgctgctgggctgttacaggaga 301 aactagagactctgttcaaagtccagggtgggggctgtgggaggaaatattagggaagcg 361 gggttcgggggataggtggtgaagctcacatccatcacgggtctctgcacacgacacagg 421 qqctccaqccaagcctgggatgtgagcacgaggctcggattgcgcatgagctctgggaaa 481 gggtgaaagcaaagacaagagttgcgggggcagggaagactgcgaggactcagggactgg 541 601 661 acggaggcgtttcggtgcgcacggagcccagccgcgttccgggaatcttgcgctcggccg 721 cgcgtggtgctcaccgcccgacccgggtgcagcgggcagctcgggtgcaggcgggggcag 781 metallothionein cap site accetetgegeceggecegetectgtgggtataatagegeteggeteetgggeteeaac bacterial cysE gene MetSerCysGluGluLeuGluIleValTrpA acqcctccaccggaccagtqqatccacaATGTCGTGTGAAGAACTGGAAATTGTCTGGA 901 snAsnIleLysAlaGluAlaArgThrLeuAlaAspCysGluProMetLeuAlaSerPheT ACAATATTAAAGCCGAAGCCAGAACGCTGGCGGACTGTGAGCCAATGCTGGCCAGTTTTT 961 yrHisAlaThrLeuLeuLysHisGluAsnLeuGlySerAlaLeuSerTyrMetLeuAlaA **ACCACGCGACGCTACTCAAGCACGAAAACCTTGGCAGTGCACTGAGCTACATGCTGGCGA** 1021 snLysLeuSerSerProIleMetProAlaIleAlaIleArgGluValValGluGluAlaT ACAAGCTGTCATCGCCAATTATGCCTGCTATTGCTATCCGTGAAGTGGTGGAAGAAGCCT 1081 yrAlaAlaAspProGluMetIleAlaSerAlaAlaCysAspIleGlnAlaValArqThrA ACGCCGCTGACCCGGAAATGATCGCCTCTGCGGCCTGTGATATTCAGGCGGTGCGTACCC 1141 rgAspProAlaValAspLysTyrSerThrProLeuLeuTyrLeuLysGlyPheHisAlaL GCGACCCGGCAGTCGATAAATACTCAACCCCGTTGTTATACCTGAAGGGTTTTCATGCCT 1201 euGlnAlaTyrArgIleGlyHisTrpLeuTrpAsnGlnGlyArgArgAlaLeuAlaIleP TGCAGGCCTATCGCATCGGTCACTGGTTGTGGAATCAGGGGCGTCGCGCACTGGCAATCT 1261

heLeuGlnAsnGlnValSerValThrPheGlnValAspIleHisProAlaAlaLysIleG

#### FIG. 2 2/3

TTCTGCAAAACCAGGTTTCTGTGACGTTCCAGGTCGATATTCACCCGGCAGCAAAAATTG

lyArgGlyIleMetLeuAspHisAlaThrGlyIleValValGlyGluThrAlaValIleG GTCGCGGTATCATGCTTGACCACGCGACAGGCATCGTCGTTGGTGAAACGGCGGTGATTG 1381

luAsnAspValSerIleLeuGlnSerValThrLeuGlyGlyThrGlyLysSerGlyGlyA AAAACGACGTATCGATTCTGCAATCTGTGACGCTTGGCGGTACGGGTAAATCTGGTGGTG 1441

spArgHisProLysIleArgGluGlyValMetIleGlyAlaGlyAlaLysIleLeuGlyAACCGTCACCCGAAAATTCGTGAAGGTGTGATGATTGGCGCGGGGCGCGAAAATCCTCGGCA1501

snlleGluValGlyArgGlyAlaLyslleGlyAlaGlySerValValLeuGlnProValP ATATTGAAGTTGGGCGCGCGCGAAGATTGGCGCAGGTTCCGTGGTGCTGCAACCGGTGC 1561

roProHisThrThrAlaAlaGlyValProAlaArgIleValGlyLysProAspSerAspLCGCCGCATACCACCGCCGCTGGCGTTCCGGCTCGTATTGTCGGTAAACCAGACAGCGATA

ysProSerMetAspMetAspGlnHisPheAsnGlyIleAsnHisThrPheGluTyrGlyA AGCCATCAATGGATATGGACCAGCATTTCAACGGTATTAACCATACATTTGAGTATGGGG 1681

spGlyIle\*\*\* growth hormone exon 5

ATGGGATCTAAtgteetgtgatetaatgteetgtgateeegetgegeettetagttgeea

gccatctgctgttacccctccctgtgccttcctagaccctggaaggtgccactccagtgc 1801

ccaccgtcctttcttaataaagcggaggaaattgcatcacattgtctgagtaggtgtcat 1861

aggggtgctgtgggctctatgggtacccaggtgctgaataattgacccggttcctcctgg 1981

ggcagaaagaagcaggcacatccccttctctgtgacacacccggtcctcgcccctggtcc 2041

ttagttccagcccactcataggacactcacagctcaggagggctccgccttcaatccca 2101

cccgctaaagtgcttggagcggtctctccctctcagccaccagccgaatctaggcctcca 2161

gagtgggaagaatttaagcaagacaggctatgaagtacagagggagagaaaatgcctcca 2221

acatgtgaggaagtgatgagagaaagcgtagaattagttttgtggcataaattttaaggt 2281

tccagctctttgtgaccccacggactgtggctgccaggctcctctgtccatgggattctc 2401

cagggcaagaatactggaggggttgccattccccaggggatcttcccagcccaaggatc 2461

aaacccgagtttctgcattgcaggcagattctttactctctgagccatcagggaagccct 2521

gtgggaaatgggaaccatgcaagaatggctttgggaccaataggaccagaatgtttggga 2581

tctgaactgggtcaagagatgtggaagagagttctaaatgcatgtgttcatgctaagtg

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FIG. 2 3/3

2641

gcttcagtcgtgtcctactatttgcaaccccgatgaactgcagccaccaggctcctctgt 2701

catgggattctccattcaagaatactggagttqagtttccttcctcccaggggatctcca 2761

aacccagggattgaccaggatctcttgtatctcctggcacttgacaggcaaatctctcac 2821

cactagcgccactggacccagtctaag--unsequenced region

61

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#### FIG. 3 1/3

#### SEQUENCE OF THE MTCK7 GENE

- 1 metallothionein promoter gaattcaaagaggaaaagtgatgaaacaaggcttggcacagactccctggtatgtaattc
- tcaggactattcaaagggaaatacccactgtcttacttcgttattggatgccagctctgc 121
- ccatcacttacaaggatgcttttcctagggggcatcctatgactagggaacctccatcct 181
- ggagccgggtggactggctaggcagtggattccctggcccattcatctattcagtcgtgg 241
- agaatgtaaggaaggctgggcgacagaaggctgagttcgctgctgggctgttacaggaga 301
- aactagagactctgttcaaagtccagggtgggggctgtggggaggaaatattagggaagcg 361
- gggttcgggggataggtggaagctcacatccatcacgggtctctgcacacgacacagg 421
- ggctccagccaagcctgggatgtgagcacgaggctcggattgcgcatgagctctgggaaa 481
- gggtgaaagcaaagacaagagttgcgggggcagggaagactgcgaggactcagggactgg 541
- tcctggagcgctacttgtcattcgggacaaagtccctccgcgttgggggggagtaggggg
- acggaggcgtttcggtgcgcacggagcccagccgcgttccgggaatcttgcgctcggccg 721
- cgcgtggtgctcaccgcccgacccgggtgcagcgggcagctcgggtgcagcgggggcag 781 metallothionein cap site \*
- accetetgegecegecegectectgtgggtataatagegeteggeteetgggeteeaac 841 bacterial cysK gene

MetSerLysIlePheGluAspAsnSer

- acgcctcccaccggaccagtggatccgtcgaccATGAGTAAGATTTTTGAAGATAACTCG 901
- LeuThrileGlyHisThrProLeuValArgLeuAsnArglleGlyAsnGlyArglleLeuCTGACTATCGGTCACACGCCGCTGGTTCGCCTGAATCGCATCGGTAACGGACGCATTCTG961
- AlaLysValGluSerArgAsnProSerPheSerValLysCysArgIleGlyAlaAsnMet GCGAAGGTGGAATCTCGTAACCCCAGCTTCAGCGTTAAGTGCCGTATCGGTGCCAACATG 1021
- TleTrpAspAlaGluLysArgGlyValLeuLysProGlyValGluLeuValGluProThr ATTTGGGATGCCGAAAAGCGCGGCGTGCTGAAACCAGGCGTTGAACTGGTTGAACCGACC 1081
- SerGlyAsnThrGlyIleAlaLeuAlaTyrValAlaAlaAlaArgGlyTyrLysLeuThr AGCGGTAATACCGGGATTGCACTGGCCTATGTAGCTGCCGCTCGCGGTTACAAACTCACC
- LeuThrMetProGluThrMetSerIleGluArgArgLysLeuLeuLysAlaLeuGlyAla CTGACCATGCCAGAAACCATGAGTATTGAACGCCGCAAGCTGCTGAAAGCGTTAGGTGCA 1201
- AsnLeuValLeuThrGluGlyAlaLysGlyMetLysGlyAlaIleGlnLysAlaGluGluAACCTGGTGCTGACGGAAGGTGCTAAAGGCATGAAAGGCGCAATCCAAAAAGCAGAAGAA

#### FIG. 3 2/3

1261

IleValAlaSerAsnProGluLysTyrLeuLeuLeuGlnGlnPheSerAsnProAlaAsn ATTGTCGCCAGCAATCCAGAGAAATACCTGCTGCTGCAACAATTCAGCAATCCGGCAAAC

ProGluIleHisGluLysThrThrGlyProGluIleTrpGluAspThrAspGlyGlnValCCTGAAATTCACGAAAAGACCACCGGTCCGGAGATATGGGAAGATACCGACGGTCAGGTT

AspValPheIleAlaGlyValGlyThrGlyGlyThrTrpThrGlyValThrProTyrIle GATGTATTTATTGCTGGCGTTGGGACTGGCGTACGTGGACTGGCGTCACGCCCTACATT

LysGlyThrLysGlyLysThrAspLeuIleSerValAlaValGluProThrAspSerPro AAAGGCACCAAAGGCAAGACCGATCTTATCTCTGTCGCCGTTGAGCCAACCGATTCTCCA 1501

VallleAlaGlnAlaLeuAlaGlyGluGluIleLysProGlyProHisLysIleGlnGlyGTTATCGCCCAGGCGCTGGCAGGTGAAGAGATTAAACCTGGCCCGCATAAAATTCAGGGT

IleGlyAlaGlyPheIleProAlaAsnLeuAspLeuLysLeuValAspLysValIleGlyATTGGCGCTGGTTTTATCCCGGCTAACCTCGATCTCAAGCTGGTCGATAAAGTCATTGGC

IleThrAsnGluGluAlaIleSerThrAlaArgArgLeuMetGluGluGluGlyIleLeu ATCACCAATGAAGAAGCGATTTCTACCGCGCGTCGTCTGATGGAAGAAGAAGGTATTCTT 1681

AlaGlyIleSerSerGlyAlaAlaValAlaAlaAlaLeuLysLeuGlnGluAspGluSer GCAGGTATCTCTTGGAGCAGCTGTTGCCGCGGCGTTGAAACTACAAGAAGATGAAAGC 1741

PheThrAsnLysAsnIleValValIleLeuProSerSerGlyGluArgTyrLeuSerThr TTTACCAACAAGAATATTGTGGTTATTCTACCATCATCGGGTGAGCGTTATTTAAGCACC 1801

AlaLeuPheAlaAspLeuPheThrGluLysGluLeuGlnGln\*\*\* growth hormone GCATTGTTTGCCGATCTCTTCACTGAGAAAGAATTGCAACAGTAAtggccagctgcgct 1861 exon 5

tctagttgccagccatctgctgttacccctccctgtgccttcctagaccctggaaggtgc 1921

cactccagtgccaccgtcctttcttaataaagcggaggaaattgcatcacattgtctga

aagacaatagcaggggtgctgtgggctctatgggtacccaggtgctgaataattgacccg 2101

gttcctcctggggcagaagcaggcacatccccttctctgtgacacacccggtcctc 2161

gcccctggtccttagttccagcccactcataggacactcacagctcaggagggctccgc 2221

cttcaatcccacccgctaaagtgcttggagcggtctctccctctcagccaccagccgaat 2281

ctaggcctccagagtgggaagaatttaagcaagacaggctatgaagtacagagggagaga 2341

aaatgcctccaacatgtgaggaagtgatgagagaaagcgtagaattagttttgtggcata 2401

actcagttqtccagctctttgtgaccccacggactgtggctgccaggctcctctgtcc

#### FIG. 3 3/3

2521

atgggattctccagggcaagaatactggaggggttgccattccccaggggatcttccca 2581

gcccaaggatcaaacccgagtttctgcattgcaggcagattctttactctctgagccatc 2641

agggaagccctgtgggaaatgggaaccatgcaagaatggctttgggaccaataggaccag 2701

aatgtttgggatctgaactgggtcaagagatgtggaagagagattctaaatgcatgtgtt 2761

catgetaagtggetteagtegtgteetaetatttgeaaceeegatgaactgeageeacea 2821

ggggatctccaaacccagggattgaccaggatctcttgtatctcctggcacttgacaggc 2941

aaatctctcaccactagcgccactggacccagtctaag---unsequenced region

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#### FIG. 4 1/5

#### SEQUENCE OF THE MTCEK1 GENE

metallothionein promoter atcatcgatcaggcagaattcaaagaggaaaagtgatgaaacaaggcttggcacagactc cctggtatgtaattctcaggactattcaaagggaaatacccactgtcttacttcgttatt 121 qqatqccaqctctqcccatcacttacaaqgatqcttttcctaqqqqqcatcctatqacta 181 gggaacctccatcctggagccgggtggactggctaggcagtggattccctggcccattca 241 tctattcagtcgtggagaatgtaaggaaggctgggcgacagaaggctgagttcgctgctg 301 ggctgttacaggagaaactagagactctgttcaaagtccagggtgggggctgtggggagga 361 421 tgcacacgacacaggggctccagccaagcctgggatgtgagcacgaggctcggattgcgc 481 atgagctctgggaaagggtgaaagcaaagacaagagttgcgggggcagggaagactgcga 541 ggactcagggactgggttcccgtaaacaccgatgactgcccacattgtggaaagctggga 601 agggggggggaatcctggagcgctacttgtcattcgggacaaagtccctccgcgttg 661 ggggcgagtaggggacggaggcgtttcggtgcgcacggagcccagccgcgttccgggaa tcttqcqctcqqccqcqtqgtqctcaccqcccqacccqggtqcaqcgggcagctcggg tqcaqqqqqqqqaaqaccetetgcqeecgqeecqcetectgtgggtataataqcqetegg bacterial cysE 841 gene metallothionein cap site -MetSerCysGluGluL

ctcctgggctccaacacgcctcccaccggaccagtggatccacaATGTCGTGTGAAGAAC

euGluIleValTrpAsnAsnIleLysAlaGluAlaArgThrLeuAlaAspCysGluProM TGGAAATTGTCTGGAACAATATTAAAGCCGAAGCCAGAACGCTGGCGGACTGTGAGCCAA 961

etLeuAlaSerPheTyrHisAlaThrLeuLeuLysHisGluAsnLeuGlySerAlaLeuS TGCTGGCCAGTTTTTACCACGCGACGCTACTCAAGCACGAAAACCTTGGCAGTGCACTGA

erTyrMetLeuAlaAsnLysLeuSerSerProIleMetProAlaIleAlaIleArgGluV GCTACATGCTGGCGAACAAGCTGTCATCGCCAATTATGCCTGCTATTGCTATCCGTGAAG

alValGluGluAlaTyrAlaAlaAspProGluMetIleAlaSerAlaAlaCysAspIleG TGGTGGAAGAAGCCTACGCCGCTGACCCGGAAATGATCGCCTCTGCGGCCTGTGATATTC

lnAlaValArgThrArgAspProAlaValAspLysTyrSerThrProLeuLeuTyrLeuL AGGCGGTGCGTACCCGCGACCCGGCAGTCGATAAATACTCAACCCCGTTGTTATACCTGA

ysGlyPheHisAlaLeuGlnAlaTyrArgIleGlyHisTrpLeuTrpAsnGlnGlyArgA AGGGTTTTCATGCCTTGCAGGCCTATCGCATCGGTCACTGGTTGTGGAATCAGGGGCGTC

#### FIG. 4 2/5

1261 rgAlaLeuAlaIlePheLeuGlnAsnGlnValSerValThrPheGlnValAspIleHisP GCGCACTGGCAATCTTTCTGCAAAACCAGGTTTCTGTGACGTTCCAGGTCGATATTCACC  ${ t rown}$  and a leading the model of the contraction of the contrac CGGCAGCAAAAATTGGTCGCGGTATCATGCTTGACCACGCGACAGGCATCGTCGTTGGTG 1381 luThrAlaVallleGluAsnAspValSerIleLeuGlnSerValThrLeuGlyGlyThrG AAACGGCGGTGATTGAAAACGACGTATCGATTCTGCAATCTGTGACGCTTGGCGGTACGG lyLysSerGlyGlyAspArgHisProLysIleArgGluGlyValMetIleGlyAlaGlyA GTAÄATCTGGTGGTGACCGTCACCCGAÄAATTCGTGAAGGTGTGATGATTGGCGCGGGCG laLysIleLeuGlyAsnIleGluValGlyArgGlyAlaLysIleGlyAlaGlySerValV CGAĀAATCCTCGGCAATATTGAAGTTGGGCGCGGGGGGGGAĀGATTGGCGCAGGTTCCGTGG alLeuGlnProValProProHisThrThrAlaAlaGlyValProAlaArgIleValGlyL TGCTGCAACCGGTGCCGCCGCATACCACCGCCGCTGGCGTTCCGGCTCGTATTGTCGGTA 1621 ysProAspSerAspLysProSerMetAspMetAspGlnHisPheAsnGlyIleAsnHisT AACCAGACAGCGATAAGCCATCAATGGATATGGACCAGCATTTCAACGGTATTAACCATA hrPheGluTyrGlyAspGlyIle\*\*\* growth hormone exon 5 CATTTGAGTÂTGGGGATGGGATCTAAtgtcctgtgatctaatgtcctgtgatcccgctgc 1741 gccttctagttgccagccatctgctgttacccctccctgtgccttcctagaccctggaag 1801 gtgccactccagtgcccaccgtcctttcttaataaagcggaggaaattgcatcacattgt 1861 1921 tgggaagacaatagcaggggtgctgtggggctctatgggtacccaggtgctgaataattga 1981 cccgqttcctcctggggcagaaagaagcaggcacatccccttctctgtgacacacccggt cctcgcccttggtccttagttccagccccactcataggacactcacagctcaggagggct ccgccttcaatcccacccgctaaagtgcttggagcggtctctccctctcagccaccagcc 2161 qaatctaggcctccagagtgggaagaatttaagcaagacaggctatgaagtacagaggga 2221 gagaaaatgcctccaacatgtgaggaagtgatgagagaaagcgtagaattagttttgtgg 2281 2341 agtcactcagttgtgtccagctctttgtgaccccacggactgtggctgccaggctcctct 2401 gtccatgggattctccagggcaagaatactggaggggttgccattccccaggggatctt 2461 2521 catcagggaagccctgtgggaaatgggaaccatgcaagaatggctttgggaccaatagga

#### FIG. 4 3/5

2581 ccagaatgtttgggatctgaactgggtcaagagatgtggaagagagattctaaatgcatg tgttcatgctaagtggcttcagtcgtgtcctactatttgcaaccccgatgaactgcaqqc metallothionein promoter 2701 atgcaagcttcagatcatcgatgaattcaaagaggaaaagtgatgaaacaaggcttggca 2761 cagactccctggtatgtaattctcaggactattcaaagggaaatacccactgtcttactt 2821 cqttattggatgccagctctgcccatcacttacaaggatgcttttcctagggggcatcct 2881 atqactagqqaacctccatcctggagccgggtggactggctaggcagtggattccctggc 2941 ccattcatctattcagtcgtggagaatgtaaggaaggctgggcgacagaaggctgagttc 3001 gctgctgggctgttacaggagaaactagagactctgttcaaagtccagggtgggggctgt 3061 gggtctctgcacacgacacaggggctccagccaagcctgggatgtgagcacgaggctcgg 3181 attgcgcatgagctctgggaaagggtgaaagcaaagacaagagttgcgggggcagggaag actgcgaggactcagggactgggttcccgtaaacaccgatgactgcccacattgtggaaa 3301 gctgggaagggcgggcaggaatcctggagcgctacttgtcattcgggacaaagtccctc 3361 cgcgttgggggggagtagggggaggaggcgtttcggtgcgcacggagcccagccgcgtt 3421 ccgggaatcttgcgctcggccgcgtggtgctcaccgcccgacccgggtgcagcgggca 3481 gctcgggtgcaggcggggcagaccctctgcgcccggcccgcctcctgtgggtataatag bacterial cysK gene 3541 metallothionein cap site MetSe cgctcggctcctgggctccaacacgcctcccaccggaccagtggatccgtcgaccATGAG rLysIlePheGluAspAsnSerLeuThrIleGlyHisThrProLeuValArgLeuAsnAr TAAGATTTTTGAAGATAACTCGCTGACTATCGGTCACACGCCGCTGGTTCGCCTGAATCG qIleGlyAsnGlyArqIleLeuAlaLysValGluSerArgAsnProSerPheSerValLy CATCGGTAACGGACGCATTCTGGCGAAGGTGGAATCTCGTAACCCCAGCTTCAGCGTTAA sCysArgIleGlyAlaAsnMetIleTrpAspAlaGluLysArgGlyValLeuLysProGl GTĞCCĞTATCGĞTĞCCAACATGATTTĞĞGATĞCCĞAAAÂĞCĞĞĞĞĞĞĞĞĞTĞCTĞAÂACCAĞĞ 3781 yValGluLeuValGluProThrSerGlyAsnThrGlyIleAlaLeuAlaTyrValAlaAl CGTTGAACTGGTTGAACCGACCAGCGGTAATACCGGGATTGCACTGGCCTATGTAGCTGC 3841

aalaargGlyTyrLysLeuThrLeuThrMetProGluThrMetSerIleGluArgArgLyCGCTCGCGGTTACAAACTCACCCTGACCATGCCAGAAACCATGAGTATTGAACGCCGCAA

#### FIG. 4 4/5

3901

sLeuLeuLysAlaLeuGlyAlaAsnLeuValLeuThrGluGlyAlaLysGlyMetLysGl GCTGCTGAAAGCGTTAGGTGCAAACCTGGTGCTGACGGAAGGTGCTAAAGGCATGAAAGG 3961

yAlaIleGlnLysAlaGluGluIleValAlaSerAsnProGluLysTyrLeuLeuLeuGl CGCAATCCAAAAAGCAGAAGAAATTGTCGCCAGCAATCCAGAGAAATACCTGCTGCTGCA 4021

nGlnPheSerAsnProAlaAsnProGluIleHisGluLysThrThrGlyProGluIleTr ACAATTCAGCAATCCGGCAAACCCTGAAATTCACGAAAAGACCACCGGTCCGGAGATATG

pGluAspThrAspGlyGlnValAspValPheIleAlaGlyValGlyThrGlyGlyThrTr GGAAGATACCGACGGTCAGGTTGATGTATTTATTGCTGGCGTTGGGACTGGCGTACGTG 4141

pThrGlyValThrProTyrIleLysGlyThrLysGlyLysThrAspLeuIleSerValAlGACTGGCGTCACGCCCTACATTAAAGGCACCAAAGGCAAGACCGATCTTATCTCTGTCGC4201

aValGluProThrAspSerProValIleAlaGlnAlaLeuAlaGlyGluGluIleLysPr CGTTGAGCCAACCGATTCTCCAGTTATCGCCCAGGCGCTGGCAGGTGAAGAGATTAAACC 4261

oGlyProHisLysIleGlnGlyIleGlyAlaGlyPheIleProAlaAsnLeuAspLeuLy TGGCCGCATAAAATTCAGGGTATTGGCGCTGGTTTTATCCCGGCTAACCTCGATCTCAA 4321

sLeuValAspLysValIleGlyIleThrAsnGluGluAlaIleSerThrAlaArgArgLe GCTGGTCGATAAAGTCATTGGCATCACCAATGAAGAAGCGATTTCTACCGCGCGTCGTCT 4381

uMetGluGluGluGlyIleLeuAlaGlyIleSerSerGlyAlaAlaValAlaAlaAlaLe GATGGAAGAAGAAGATTCTTGCAGGTATCTCTTGGAGCAGCTGTTGCCGCGGCGTT

uLysLeuGlnGluAspGluSerPheThrAsnLysAsnIleValValIleLeuProSerSe GAAACTACAAGAAGATGAAAGCTTTACCAACAAGAATATTGTGGTTATTCTACCATCATC 4501

nGln\*\*\* growth hormone exon 5

cttcctagaccctggaaggtgccactccagtgcccaccgtcctttcttaataaagcggag 4681

gaaattgcatcacattgtctgagtaggtgtcattctattctagggggtggggtcgggcag 4741

gatagegaggggggggattgggaagaeaatageaggggtgetgtggggetetatgggtaee

ctctgtgacacacccggtcctcgcccctggtccttagttccagccccactcataggacac 4921

tcacagetcaggagggctccgcettcaatcccacccgctaaagtgcttggagcggtctct 4981

ccctctcaqccaccaqccgaatctaggcctccagagtgggaagaatttaagcaagacagg

# FIG. 4 5/5

ccccgatgaactgcaggcatgcaagcttcagctgc

# FIG. 5 1/3

# SEQUENCE OF THE MTACEA2 GENE

1 metallothionein promoter
gaattcaaagaggaaaagtgatgaaacaaggcttggcacagactccctggtatgtaattc 61
tcaggactattcaaagggaaatacccactgtcttacttcgttattggatgccagctctgc 121
ccatcacttacaaggatgcttttcctagggggcatcctatgactagggaacctccatcct 181
ggagccgggtggactggctaggcagtggattccctggcccattcatctattcagtcgtgg 241
agaatgtaaggaaggctgggcgacagaaggctgagttcgctgctgggctgttacaggaga 301
aactagagactctgttcaaagtccagggtgggggctgtggggaggaaatattagggaagcg 361
gggttcgggggataggtggagctcacatccatcacgggtctctgcacacgacacagg 421
ggctccagccaagcctgggatgtgagcacgaggctcggattgcgcatgagctctgggaaa 481
gggtgaaagcaaagactgcgggggcagggaagactgcgaggactcagggactgg 541
gttcccgtaaacaccgatgactgcccacattgtggaaagctgggaagggcgggc
tcctggagcgctacttgtcattcgggacaaagtccctccgcgttgggggggagtaggggg
acggaggcgtttcggtgcgcacggagcccagccgcgttccgggaatcttgcgctcggccg 721
cgcgtggtgctcaccgcccgacccgggtgcagcggggcagctcgggtgcaggcgggggcag 781 metallothionein cap site *
accetetgegeeegeeteetgtgggtataatagegeteggeteetgggeteeaac 841 bacterial <u>ace A</u> sequence
MetLysThrArgThrGlnG acgcctccaccggaccagtggatcctctagagtcatcaccATGAAAACCCGTACACAAC
901 lnIleGluGluLeuGlnLysGluTrpThrGlnProArgTrpGluGlyIleThrArgProT
AAATTGAAGAATTACAGAAAGAGTGGACTCAACCGCGTTGGGAAGGCATTACTCGCCCAT
yrSerAlaGluAspValValLysLeuArgGlySerValAsnProGluCysThrLeuAlaG ACAGTGCGGAAGATGTGGTGAAATTACGCGGTTCAGTCAATCCTGAATGCACGCTGGCGC
1021 lnLeuGlyAlaAlaLysMetTrpArgLeuLeuHisGlyGluSerLysLysGlyTyrIleA
AACTGGGCGCAGCGAAAATGTGGCGTCTGCTGCACGGTGAGTCGAAAAAAGGCTACATCA
snSerLeuGlyAlaLeuThrGlyGlyGlnAlaLeuGlnGlnAlaLysAlaGlyIleGluAACAGCCTCGGCGCACTGACTGGCGGTCAGGCGCTGCAACAGGCGAAAGCGGGTATTGAAG
1141
lavalTyrLeuSerGlyTrpGlnValAlaAlaAspAlaAsnLeuAlaAlaSerMetTyrPCAGTCTATCTGTCGGGATGGCAGGTAGCGGCGGACGCTAACCTGGCGGCCAGCATGTATC
roAspGlnSerLeuTyrProAlaAsnSerValProAlaValValGluArgIleAsnAsnT CGGATCAGTCGCTCTATCCGGCAAACTCGGTGCCAGCTGTGGTGGAGCGGATCAACAACA

#### FIG. 5 2/3

1261 hrPheArgArgAlaAspGlnIleGlnTrpSerAlaGlyIleGluProGlyAspProArgT CCTTCCGTCGTGCCGATCAGATCCAATGGTCCGCGGGCATTGAGCCGGGCGATCCGCGCT yrValAspTyrPheLeuProIleValAlaAspAlaGluAlaGlyPheGlyGlyValLeuA **ĀTGTCGĀTTĀCTTCCTGCCGATCGTTGCCGATGCGGAAGCCGGTTTTTGGCGGTGTCCTGA** snAlaPheGluLeuMetLysAlaMetIleGluAlaGlyAlaAlaAlaValHisPheGluA **ATGCCTTTGAACTGATGAĀAGCGATGATTGAAGCCGGTGCAGCGGCAGTTCACTTCGAAG** spGlnLeuAlaSerValLysLysCysGlyHisMetGlyGlyLysValLeuValProThrG **ATCAGCTGGCGTCAGTGAĀGAĀATĞCGGTCACATGGGCGGCAĀAGTTTTAGTGCCAACTC** lnGluAlaIleGlnLysLeuValAlaAlaArgLeuAlaAlaAspValThrGlyValProT hrLeuLeuValAlaArgThrAspAlaAspAlaAlaAspLeuIleThrSerAspCysAspP CCCTGCTGGTTGCCCGTACCGATGCTGATGCGGCGGATCTGATCACCTCCGATTGCGACC roTyrAspSerGluPheIleThrGlyGluArgThrSerGluGlyPhePheArgThrHisA CGTATGACAGCGAATTTATTACCGGCGAGCGTACCAGTGAAGGCTTCTTCCGTACTCATG 1681 laGlyIleGluGlnAlaIleSerArgGlyLeuAlaTyrAlaProTyrAlaAspLeuValT CGGGCATTGAGCAAGCGATCAGCCGTGGCCTGGCGTATGCGCCATATGCTGACCTGGTCT 1741 rpCysGluThrSerThrProAspLeuGluLeuAlaArgArgPheAlaGlnAlaIleHisA GGTGTGAAACCTCCACGCCGGATCTGGAACTGGCGCGTCGCTTTGCACAAGCTATCCACG laLysTyrProGlyLysLeuLeuAlaTyrAsnCysSerProSerPheAsnTrpGlnLysA CGAĀATĀTCCGGGCĀĀACTGCTGGCTTĀTAACTGCTCGCCGTCGTTCAACTGGCAGAĀAA snLeuAspAspLysThrIleAlaSerPheGlnGlnGlnLeuSerAspMetGlyTyrLysP ACCTCGACGACAAAACTATTGCCAGCTTCCAGCAGCAGCTGTCGGATATGGGCTACAAGT heGlnPheIleThrLeuAlaGlyIleHisSerMetTrpPheAsnMetPheAspLeuAlaA TCCAGTTCATCACCCTGGCAGGTATCCACAGCATGTGGTTCAACATGTTTGACCTGGCAA snAlaTyrAlaGlnGlyGluGlyMetLysHisTyrValGluLysValGlnGlnProGluP ACGCCTATGCCCAGGGCGAGGGTATGAAGCACTACGTTGAGAAAGTGCAGCAGCCGGAAT heAlaAlaAlaLysAspGlyTyrThrPheValSerHisGlnGlnGluValGlyThrGlyT TTGCCGCCGCGAAAGATGGCTATACCTTCGTATCTCACCAGCAGGAAGTGGGTACAGGTT 2101 yrPheAspLysValThrThrIleIleGlnGlyGlyAspValPheSerHisArgAlaAspA ACTTCGATAAAGTGACGACTATTATTCAGGGCGGCGACGTCTTCAGTCACCGCGCTGACC growth hormone exon 5 2161 rqLeuHis\*\*\* ctgttacccctccctgtgccttcctagaccctggaaggtgccactccagtgcccaccgtc 2281 ctttcttaataaagcggaggaaattgcatcacattgtctgagtaggtgtcattctattct

#### FIG. 5 3/3

2341

agggggtggggtcgggcaggatagcgaggggggggattgggaagacaatagcaggggtgc

tgtgggctctatgggtacccaggtgctgaataattgacccggttcctcctggggcagaaa 2461

gaagcaggcacatccccttctctgtgacacacccggtcctcgccctggtccttagttcc 2521

agecceacteataggacacteacageteaggagggeteegeetteaateccaccegetaa 2581

agtgcttggagcggtctctccctctcagccaccagccgaatctaggcctccagagtggga 2641

agaatttaagcaagacaggctatgaagtacagagggagagaaaatgcctccaacatgtga 2701

ggaagtgatgagagaaagcgtagaattagttttgtggcataaattttaaggtgactacac 2761

acttggcccaactacccttgggaaatgtgtgtgtgttagtcactcagttgtgtccagctc 2821

tttgtgacccacggactgtggctgccaggctcctctgtccatgggattctccagggcaa 2881

gaatactggaggggttgccattccccaggggatcttcccagcccaaggatcaaacccga 2941

gtttctgcattgcaggcagattctttactctctgagccatcagggaagccctgtgggaaa 3001

tgggaaccatgcaagaatggctttgggaccaataggaccagaatgtttgggatctgaact 3061

gggtcaagagatgtggaagagattctaaatgcatgtgttcatgctaagtggcttcagt 3121

cgtgtcctactatttgcaaccccgatgaactgcag

#### FIG. 6 1/3

#### SEQUENCE OF THE MTAceB2 GENE

- 1 metallothionein promoter gaattcaaagaggaaaagtgatgaaacaaggcttggcacagactccctggtatgtaattc 61 tcaggactattcaaagggaaatacccactgtcttacttcgttattggatgccagctctgc 121 ccatcacttacaaggatgcttttcctagggggcatcctatgactagggaacctccatcct 181  $\tt ggagccgggtggactggctaggcagtggattccctggcccattcatctattcagtcgtgg$ 241 agaatgtaaggaaggctgggcgacagaaggctgagttcgctgctgggctgttacaggaga 301 aactagagactctgttcaaagtccagggtgggggctgtggggggaaatattagggaagcg 361 gggttcgggggataggtggaagctcacatccatcacgggtctctgcacacgacacagg 421 ggctccagccaagcctgggatgtgagcacgaggctcggattgcgcatgagctctqggaaa 481 gggtgaaagcaaagacaagagttgcgggggcagggaagactgcgaggactcagggactgg 541 601 tcctggagcgctacttgtcattcgggacaaagtccctccgcgttgggggcgagtaggggg 661 acggaggcgtttcggtgcgcacggagcccagccgcgttccgggaatcttgcgctcggccg 721 cgcgtggtgctcaccgcccgacccgggtgcagctgggcagctcgggtgcagggggggag metallothionein cap site 781 accetetgegeeeggeeegeeteetgtgggtataatagegeteggeteetgggeteeaae bacterial aceB sequence 841 MetThrGluGlnAlaThrT acgcctcccaccggaccagtggatcctctagagtcatcaccATGACTGAACAGGCAACAA 901 hrThrAspGluLeuAlaPheThrArgProTyrGlyGluGlnGluLysGlnIleLeuThrA CAACCGATGAACTGGCTTTCACAAGGCCGTATGGCGAGCAGGAGAAGCAAATTCTTACTG 961 laGluAlaValGluPheLeuThrGluLeuValThrHisPheThrProGlnArgAsnLysL CCGAAGCGGTAGAATTTCTGACTGAGCTGGTGACGCATTTTACGCCACAACGČAATAÃAC 1021
- euLeuAlaAlaArgIleGlnGlnGlnGlnAspIleAspAsnGlyThrLeuProAspPheI TTCTGGCAGCGCGCATTCAGCAGCAGCAAGATATTGATAACGGAACGTTGCCTGATTTTA
- leSerGluThrAlaSerIleArgAspAlaAspTrpLysIleArgGlyIleProAlaAspL TTTCGGAAACAGCTTCCATTCGCGATGCTGATTGGAAAATTCGCGGGATTCCTGCGGACT
- euGluAspArgArgValGluIleThrGlyProValGluArgLysMetValIleAsnAlaL TAGAAGACCGCCGCTAGAGATAACTGGCCCGGTAGAGCGCAAGATGGTGATCAACGCGC 1201
- euAsnAlaAsnValLysValPheMetAlaAspPheGluAspSerLeuAlaProAspTrpA TCAACGCCAATGTGAAAGTCTTTATGGCCGATTTCGAAGATTCACTGGCACCAGACTGGA

# 18/25 FIG. 6 2/3

- 1261
  snLysValileAspGlyGlnileAsnLeuArgAspAlaValAsnGlyThrileSerTyrT
  ACAAAGTGATCGACGGGCAAATTAACCTGCGTGATGCGGTTAACGGCACCATCAGTTACA
  1321
- hrAsnGluAlaGlyLysIleTyrGlnLeuLysProAsnProAlaValLeuIleCysArgV CCAATGAAGCAGGCAAAATTTACCAGCTCAAGCCCAATCCAGCGGTTTTGATTTGTCGGG
- alargGlyLeuHisLeuProGluLysHisValThrTrpArgGlyGluAlaIleProGlyS TACGCGGTCTGCACTTGCCGGAAAAACATGTCACCTGGCGTGATGAGGCAATCCCCGGCA
- erLeuPheAspPheAlaLeuTyrPhePheHisAsnTyrGlnAlaLeuLeuAlaLysGlyS GCCTGTTTGATTTTGCGCTCTATTTCTTCCACAACTATCAGGCACTGTTGGCAAAGGGCA 1501
- erGlyProTyrPheTyrLeuProLysThrGlnSerTrpGlnGluAlaAlaTrpTrpSerG GTGGTCCCTATTTCTATCTGCCGAAAACCCAGTCCTGGCAGGAAGCGGCCTGGTGGAGCG
- luValPheSerTyrAlaGluAspArgPheAsnLeuProArgGlyThrIleLysAlaThrL AAGTCTTCAGCTATGCAGAAGATCGCTTTAATCTGCCGCGCGCACCATCAAGGCGACGT 1621
- euLeuIleGluThrLeuProAlaValPheGlnMetAspGluIleLeuHisAlaLeuArgA TGCTGATTGAAACGCTGCCCGCCGTGTTCCAGATGGATGAAATCCTTCACGCGCTGCGTG
- sphisllevalGlyLeuAsnCysGlyArgTrpAspTyrllePheSerTyrlleLysThrL ACCATATTGTTGGTCTGAACTGCGGTCGTTGGGATTACATCTTCAGCTATATCAAAACGT 1741
- euLysAsnTyrProAspArgValLeuProAspArgGlnAlaValThrMetAspLysProP TGAAAAACTATCCCGATCGCGTCCTGCCAGACAGACAGGCAGTGACGATGGATAAACCAT
- heLeuAsnAlaTyrSerArgLeuLeuIleLysThrCysHisLysArgGlyAlaPheAlaM TCCTGAATGCTTACTCACGCCTGTTGATTAAAACCTGCCATAAACGCGGTGCTTTTGCGA 1861
- etGlyGlyMetAlaAlaPhelleProSerLysAspGluGluHisAsnAsnGlnValLeuA TGGGCGGCATGGCGGCGTTTATTCCGAGCAAAGATGAAGAGCACAATAACCAGGTGCTCA
- snLysVallysAlaAspLysSerLeuGluAlaAsnAsnGlyHisAspGlyThrTrpIleA ACAAAGTAAAAGCGGATAAATCGCTGGAAGCCAATAACGGTCACGATGGCACATGGATCG
- laHisProGlyLeuAlaAspThrAlaMetAlaValPheAsnAspIleLeuGlySerArgL CTCACCCAGGCCTTGCGGACACGGCAATGGCGGTATTCAACGACATTCTCGGCTCCCGTA 2041
- ysAsnGlnLeuGluValMetArgGluGlnAspAlaProIleThrAlaAspGlnLeuLeuA AAAATCAGCTTGAAGTGATGCGCGAACAAGACGCGCCGATTACTGCCGATCAGCTGCTGG 2101
- laproCysAspGlyGluArgThrGluGluGlyMetArgAlaAsnIleArgValAlaValG CACCTTGTGATGGTGAACGCACCGAAGAAGGTATGCGCGCCAACATTCGCGTGGCTGTGC 2161
- lnTyrIleGluAlaTrpIleSerGlyAsnGlyCysValProIleTyrGlyLeuMetGluA AGTACATCGAAGCGTGGATCTCTGGCAACGGCTGTGTGCCGATTTATGGCCTGATGGAAG 2221
- spalaAlaThrAlaGluIleSerArgThrSerIleTrpGlnTrpIleHisHisGlnLysT ATGCGGCGACGGCTGAAATTTCCCGTACCTCGATCTGGCAGTGGATCCATCATCAAAAAA

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## FIG. $6 \ 3/3$

2281 hrLeuSerAsnGlyLysProValThrLysAlaLeuPheArgGlnMetLeuGlyGluGluM CGTTGAGCAATGGCAAACCGGTGACCAAAGCCTTGTTCCGCCAGATGCTGGGCGAAGAGA etLysValIleAlaSerGluLeuGlyGluGluArgPheSerGlnGlyArgPheAspAspA TGAÃAGTCATTGCCAGCGAACTGGGCGAAGAACGTTTCTCCCAGGGGCGTTTTGACGATG 2401 laAlaArgLeuMetGluGlnIleThrThrSerAspGluLeuIleAspPheLeuThrLeuP CCGCACGCTTGATGGAACAGATCACCACTTCCGATGAGTTAATTGATTTCCTGACCCTGC growth hormone exon 5 roGlyTyrArgLeuLeuAla\*\*\* CAGGCTACCGCCTGTTAGCGTAAtttgacctgcgccttctagttgccagccatctgctqt tacccctccctqtqccttcctagaccctggaaggtgccactccagtgcccaccgtccttt 2581 cttaataaagcggaggaaattgcatcacattgtctgagtaggtgtcattctattctagqq 2641 ggtggggtcgggcaggatagcgagggggaggattgggaagacaatagcaggggtgctgtg 2701 2761 caggcacatccccttctctgtgacacacccggtcctcgccctggtccttagttccagcc 2821 ccactcataggacactcacagctcaggagggctccgccttcaatcccacccgctaaagtg cttggagcggtctctccctctcagccaccagccgaatctaggcctccagagtgggaagaa tttaagcaagacaggctatgaagtacagagggagagaaaatgcctccaacatgtgaggaa gtgatgagagaaagcgtagaattagttttgtggcataaattttaaggtgactacacactt 3061 ggcccaactacccttgggaaatgtgtgtgtgttagtcactcagttgtgtccagctctttg 3121 tgaccccacggactgtggctgccaggctcctctgtccatgggattctccagggcaagaat 3181 actggaggggttgccattccccaggggatcttcccagcccaaggatcaaacccgagttt 3241 ctgcattgcaggcagattctttactctctgagccatcagggaagccctgtgggaaatggg 3301 aaccatgcaagaatggctttgggaccaataggaccagaatgtttgggatctgaactgggt 3361 caagagatgtggaagagattctaaatgcatgtgttcatgctaagtggcttcagtcgtg 3421 tcctactatttgcaaccccgatgaactgcag

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# FIG. 7 1/5

# SEQUENCE OF THE MTACEAB1 GENE

1 metallothionein promoter gaattcaaagaggaaaagtgatgaaacaaggcttggcacagactccctggtatgtaattc 61 tcaggactattcaaagggaaatacccactgtcttacttcgttattggatgccagctctqc 121 ccatcacttacaaggatgcttttcctagggggcatcctatgactagggaacctccatcct 181 ggagccgggtggactggctaggcagtggattccctggcccattcatctattcagtcgtgg 241 agaatgtaaggaaggctgggcgacagaaggctgagttcgctgctgggctgttacaggaga 301 aactagagactctgttcaaagtccagggtgggggctgtggggaggaaatattagggaagcg 361 gggttcgggggataggtggaagctcacatccatcacgggtctctgcacacgacacagg 421 ggctccagccaagcctgggatgtgagcacgaggctcggattgcgcatgagctctgggaaa 481 gggtgaaagcaaagacaagagttgcgggggcagggaagactgcgaggactcagggactgg 541 601 661 acggaggcgtttcggtgcgcacggagcccagccgcgttccgggaatcttgcgctcggccg 721 cgcgtggtgctcaccgcccgacccgggtgcagcggggcagctcgggtgcaggcgggggcag metallothionein cap site 781 accetetgegeeeggeeegeeteetgtgggtataatagegeteggeteetgggeteeaae bacterial aceA sequence MetLysThrArgThrGlnG acqcctcccaccggaccagtggatcctctagagtcatcaccATGAAAACCCGTACACAAC 901 lnIleGluGluLeuGlnLysGluTrpThrGlnProArgTrpGluGlyIleThrArgProT AAATTGAAGAATTACAGAAAGAGTGGACTCAACCGCGTTGGGAAGGCATTACTCGCCCAT yrSerAlaGluAspValValLysLeuArgGlySerValAsnProGluCysThrLeuAlaG ACAGTGCGGAAGATGTGGTGAAATTACGCGGTTCAGTCAATCCTGAATGCACGCTGGCGC 1021 lnLeuGlyAlaAlaLysMetTrpArgLeuLeuHisGlyGluSerLysLysGlyTyrIleA AACTGGGCGCAGCGAAAATGTGGCGTCTGCTGCACGGTGAGTCGAAAAAAGGCTACATCA snSerLeuGlyAlaLeuThrGlyGlyGlnAlaLeuGlnGlnAlaLysAlaGlyIleGluA ACAGCCTCGGCGCACTGACTGGCGGTCAGGCGCTGCAACAGGCGAAAGCGGGTATTGAAG 1141 laValTyrLeuSerGlyTrpGlnValAlaAlaAspAlaAsnLeuAlaAlaSerMetTyrP CAGTCTATCTGTCGGGATGGCAGGTAGCGGCGGACGCTAACCTGGCGGCCAGCATGTATC

roAspGlnSerLeuTyrProAlaAsnSerValProAlaValValGluArgIleAsnAsnTCGGATCAGTCGCTCTATCCGGCAAACTCGGTGCCAGCTGTGGTGGAGCGGATCAACAACA

1201

2281

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#### FIG. 7 2/5

1261 hrPheArgArgAlaAspGlnIleGlnTrpSerAlaGlyIleGluProGlyAspProArgT CCTTCCGTCGTGCCGATCAGATCCAATGGTCCGCGGCCATTGAGCCGGGCGATCCGCGCT 1321 yrValAspTyrPheLeuProIleValAlaAspAlaGluAlaGlyPheGlyGlyValLeuA **ĀTGTCGĀĪTĀCTTCCTGCCGATCGTTGCCGATGCGGAAGCCGGTTTTGGCGGTGTCCTGA** 1381 snAlaPheGluLeuMetLysAlaMetIleGluAlaGlyAlaAlaAlaValHisPheGluA ATGCCTTTGAACTGATGAĀAGCGATGATTGAAGCCGGTGCAGCGGCAGTTCACTTCGAAG spGlnLeuAlaSerValLysLysCysGlyHisMetGlyGlyLysValLeuValProThrG **ATCAGCTGGCGTCAGTGAĀGAĀATĞCGGTCACATGGGCGGCAĀAGTTTTAGTGCCAACTC** 1501 lnGluAlaIleGlnLysLeuValAlaAlaArgLeuAlaAlaAspValThrGlyValProT hrLeuLeuValAlaArgThrAspAlaAspAlaAlaAspLeuIleThrSerAspCysAspP CCCTGCTGGTTGCCCGTACCGATGCTGATGCGGCGGATCTGATCACCTCCGATTGCGACC roTyrAspSerGluPheIleThrGlyGluArgThrSerGluGlyPhePheArgThrHisA CGTATGACAGCGAATTTATTACCGGCGAGCGTACCAGTGAAGGCTTCTTCCGTACTCATG 1681 laGlyIleGluGlnAlaIleSerArgGlyLeuAlaTyrAlaProTyrAlaAspLeuValT CGGGCATTGAGCAAGCGATCAGCCGTGGCCTGGCGTATGCGCCATATGCTGACCTGGTCT 1741 rpCysGluThrSerThrProAspLeuGluLeuAlaArgArgPheAlaGlnAlaIleHisA GGTGTGAAACCTCCACGCCGGATCTGGAACTGGCGCGTCGCTTTGCACAAGCTATCCACG laLysTyrProGlyLysLeuLeuAlaTyrAsnCysSerProSerPheAsnTrpGlnLysA CGAÃATÃTCCGGGĈAÃACTGCTGGCTTĀTAACTĞCTCGCCGTCGTTCAACTGGCAGAAAA 1861  $\verb|snLeuAspAspLysThrIleAlaSerPheGlnGlnGlnLeuSerAspMetGlyTyrLysP|\\$ ACCTCGACGACAAAACTATTGCCAGCTTCCAGCAGCAGCTGTCGGATATGGGCTACAAGT heGlnPheIleThrLeuAlaGlyIleHisSerMetTrpPheAsnMetPheAspLeuAlaA TCCAGTTCATCACCCTGGCAGGTATCCACAGCATGTGGTTCAACATGTTTGACCTGGCAA 1981  $\verb|snAlaTyrAlaGlnGlyGluGlyMetLysHisTyrValGluLysValGlnGlnProGluP|\\$ ACGCCTATGCCCAGGGCGAGGGTATGAAGCACTACGTTGAGAAAGTGCAGCAGCCGGAAT 2041  $\verb|heAlaAlaAlaLysAspGlyTyrThrPheValSerHisGlnGlnGluValGlyThrGlyT|\\$ TTGCCGCCGCGAAAGATGGCTATACCTTCGTATCTCACCAGCAGGAAGTGGGTACAGGTT 2101 yrPheAspLysValThrThrIleIleGlnGlyGlyAspValPheSerHisArgAlaAspA ACTTCGATAAAGTGACGACTATTATTCAGGGCGGCGACGTCTTCAGTCACCGCGCTGACC growth hormone exon 5 2161 rgLeuHis\*\*\* 2221

ctgttacccctccctgtgccttcctagaccctggaaggtgccactccagtgcccaccgtc

 $\verb|cttcttaataaagcggaggaaattgcatcacattgtctgagtaggtgtcattctattct|\\$ 

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# 22/25 FIG. 7 3/5

2341 agggggtggggtcgggcaggatagcgagggggaggattgggaagacaatagcaggggtgc 2401 tgtgggctctatgggtacccaggtgctgaataattgacccggttcctcctggggcagaaa 2461 gaagcaggcacatccccttctctgtgacacacccggtcctcgcccctggtccttagttcc 2521 agccccactcataggacactcacagctcaggagggctccgccttcaatcccacccgctaa 2581 agtgcttggagcggtctctccctctcagccaccagccgaatctaggcctccagagtggga 2641 agaatttaagcaagacaggctatgaagtacagagggagagaaaatgcctccaacatgtga 2701 ggaagtgatgagagaaagcgtagaattagttttgtggcataaattttaaggtgactacac 2761 acttggcccaactacccttgggaaatgtgtgtgtgttagtcactcagttgtgtccagctc 2821 tttqtqaccccacqqactqtqqctqccaqqctcctctqtccatqqqattctccaqqqcaa 2881 gaatactggagggggttgccattccccaggggatcttcccagcccaaggatcaaacccga 2941 qtttctgcattgcaggcagattctttactctctgagccatcagggaagccctgtgggaaa 3001 tgggaaccatgcaagaatggctttgggaccaataggaccagaatgtttgggatctgaact 3061 gggtcaagagatgtggaagagattctaaatgcatgtgttcatgctaagtggcttcagt metallothionein promoter 3121 cgtgtcctactatttgcaaccccgatgaactgcaggaattcaaagaggaaaagtgatgaa 3181 acaaggcttggcacagactccctggtatgtaattctcaggactattcaaagggaaatacc cactgtcttacttcgttattggatgccagctctgcccatcacttacaaggatgcttttcc tagggggcatcctatgactagggaacctccatcctggagccgggtggactggctaggcag gaaggctgagttcgctgctgggctgttacaggagaaactagagactctgttcaaagtcca qqqtqqqqctqtqqqaqqaaatattaqqqaaqcqqqqttcqqqqqataqqtqqtqaaqc 3541 tcacatccatcacgggtctctgcacacgacacaggggctccagccaagcctgggatgtga 3601 gcacgaggctcggattgcgcatgagctctgggaaagggtgaaagcaaagacaagagttgc 3661 gggggcagggaagactgcgaggactcagggactgggttcccgtaaacaccgatgactgcc 3721 cacattgtggaaagctgggaagggcgggcaggaatcctggagcgctacttgtcattcgg 3781 gacaaagtccctccgcgttggggcgagtaggggggacggaggcgtttcggtgcgcacgga WO 92/18635 PCT/AU92/00164

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## FIG. 7 4/5

3841 gcccagccgcgttccgggaatcttgcgctcggccgcgcgtggtgctcaccgcccgacccg 3901 ggtgcagcgggcagctcgggtgcaggcggggcagaccctctgcgcccggcccgcctcct 3961 metallothionein cap site gtgggtataatagcgctcggctcctgggctccaacacgcctcccaccggaccagtggatc bacterial aceB sequence 4021 MetThrGluGlnAlaThrThrThrAspGluLeuAlaPheThrAr ctctagagtcatcaccATGACTGAACAGGCAACAACCAGTGAACTGGCTTTCACAAG  ${\tt gProTyrGlyGluGlnGluLysGlnIleLeuThrAlaGluAlaValGluPheLeuThrGlu$ 4141 uLeuValThrHisPheThrProGlnArgAsnLysLeuLeuAlaAlaArgIleGlnGlnGl GCTGGTGACGCATTTTACGCCACAACGCAATAAACTTCTGGCAGCGCGCATTCAGCAGCA nGlnAspIleAspAsnGlyThrLeuProAspPheIleSerGluThrAlaSerIleArgAs GCAAGATATTGATAACGGAACGTTGCCTGATTTTATTTCGGAAACAGCTTCCATTCGCGA pAlaAspTrpLysIleArgGlyIleProAlaAspLeuGluAspArgArgValGluIleTh TGCTGATTGGAAAATTCGCGGGATTCCTGCGGACTTAGAAGACCGCCGCGTAGAGATAAC 4321  $\verb"rGlyProValGluArgLysMetValIleAsnAlaLeuAsnAlaAsnValLysValPheMe"$ TGGCCCGGTAGAGCGCAAGATGGTGATCAACGCGCTCAACGCCAATGTGAAAGTCTTTAT 4381 tAlaAspPheGluAspSerLeuAlaProAspTrpAsnLysValIleAspGlyGlnIleAs GGCCGATTTCGAAGATTCACTGGCACCAGACTGGAACAAAGTGATCGACGGGCAAATTAA nLeuArgAspAlaValAsnGlyThrIleSerTyrThrAsnGluAlaGlyLysIleTyrGl CCTGCGTGATGCGGTTAACGGCACCATCAGTTACACCAATGAAGCAGGCAAAATTTACCA 4501  ${ t nLeuLysProAsnProAlaValLeuIleCysArgValArgGlyLeuHisLeuProGluLy}$ GCTCAAGCCCAATCCAGCGGTTTTGATTTGTCGGGTACGCGGTCTGCACTTGCCGGAAAÂ sHisValThrTrpArgGlyGluAlaIleProGlySerLeuPheAspPheAlaLeuTyrPh **ACATGTCACCTGGCGTGGTGAGGCAATCCCCGGCAGCCTGTTTGATTTTGCGCTCTATTT** 4621 ePheHisAsnTyrGlnAlaLeuLeuAlaLysGlySerGlyProTyrPheTyrLeuProLy CTTCCACAACTÂTCAGGCACTGTTGGCAAÂGGGCAGTGGTCCCTÂTTTCTÂTCTGCCGAÂ sThrGlnSerTrpGlnGluAlaAlaTrpTrpSerGluValPheSerTyrAlaGluAspAr **AACCCAGTCCTGGCAGGAAGCGGCCTGGTGGAGCGAAGTCTTCAGCTATGCAGAAGA**TCG 4741 gPheAsnLeuProArgGlyThrIleLysAlaThrLeuLeuIleGluThrLeuProAlaVa CTTTAATCTGCCGCGCGCACCATCAAGGCGACGTTGCTGATTGAAACGCTGCCCGCCGT 4801 lPheGlnMetAspGluIleLeuHisAlaLeuArgAspHisIleValGlyLeuAsnCysGl GTTCCAGATGGATGAAATCCTTCACGCGCTGCGTGACCATATTGTTGGTCTGAACTGCGG 4861 yArgTrpAspTyrIlePheSerTyrIleLysThrLeuLysAsnTyrProAspArgValLe TCGTTGGGATTACATCTTCAGCTATATCAAAACGTTGAAAAACTATCCCGATCGCGTCCT

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4921 uProAspArgGlnAlaValThrMetAspLysProPheLeuAsnAlaTyrSerArgLeuLe GCCAGACAGACAGGCAGTGACGATGGATAÃACCATTCCTGAATGCTTÃCTCACGCCTGTT ulleLysThrCysHisLysArgGlyAlaPheAlaMetGlyGlyMetAlaAlaPheIlePr GATTAAAACCTGCCATAAACGCGGTGCTTTTGCGATGGGCGGCATGGCGGCGTTTATTCC oSerLysAspGluGluHisAsnAsnGlnValLeuAsnLysValLysAlaAspLysSerLe GAGCAĀAGAĪGAAGAGCACAATAACCAGGTGCTCAACAĀAGTAAĀAGCGGAĪAĀATCGCT uGluAlaAsnAsnGlyHisAspGlyThrTrpIleAlaHisProGlyLeuAlaAspThrAl GGAAGCCAATAACGGTCACGATGGCACATGGATCGCTCACCCAGGCCTTGCGGACACGGC aMetAlaValPheAsnAspIleLeuGlySerArgLysAsnGlnLeuGluValMetArgGl AATGGCGGTATTCAACGACATTCTCGGCTCCCGTAAAAATCAGCTTGAAGTGATGCGCGA uGlnAspAlaProIleThrAlaAspGlnLeuLeuAlaProCysAspGlyGluArgThrGl ACAAGACGCGCCGATTACTGCCGATCAGCTGCTGGCACCTTGTGATGGTGAACGCACCGA 5281 uGluGlyMetArgAlaAsnIleArgValAlaValGlnTyrIleGluAlaTrpIleSerGl AGAAGGTATGCGCGCCAACATTCGCGTGGCTGTGCAGTĀCATCGAAGCGTGGATCTCTGG 5341 yAsnGlyCysValProIleTyrGlyLeuMetGluAspAlaAlaThrAlaGluIleSerAr CAACGGCTGTGTGCCGATTTATGGCCTGATGGAAGATGCGGCGACGGCTGAAATTTCCCG 5401 gThrSerIleTrpGlnTrpIleHisHisGlnLysThrLeuSerAsnGlyLysProValTh TACCTCGATCTGGCAGTGGATCCATCATCAAAAAACGTTGAGCAATGGCAAACCGGTGAC rLysAlaLeuPheArgGlnMetLeuGlyGluGluMetLysValIleAlaSerGluLeuGl CAÂAGCCTTGTTCCGCCAGATGCTGGGCGAAGAGATGAÂAGTCATTGCCAGCGAACTGGG 5521 yGluGluArgPheSerGlnGlyArgPheAspAspAlaAlaArgLeuMetGluGlnIleTh <u>CGAAGAACGTTTCTCCCAGGGGCGTTTTGACGATGCCGCACGCTTGATGGAACAGATCAC</u> 5581 rThrSerAspGluLeuIleAspPheLeuThrLeuProGlyTyrArgLeuLeuAla\*\*\* CACTTCCGATGAGTTAATTGATTTCCTGACCCTGCCAGGCTACCGCCTGTTAGCGTAAtt growth hormone exon 5 5641 5701 cctggaaggtgccactccagtgcccaccgtcctttcttaataaagcggaggaaattgcat cacattgtctgagtaggtgtcattctattctagggggtggggtcgggcaggatagcgagg gggaggattgggaagacaatagcaggggtgctgtggggctctatgggtacccaggtgctga 5881 ataattgacccggttcctcctggggcagaaagaagcaggcacatccccttctctgtgaca

cacccggtcctcgcccttggtccttagttccagccccactcataggacactcacagctca

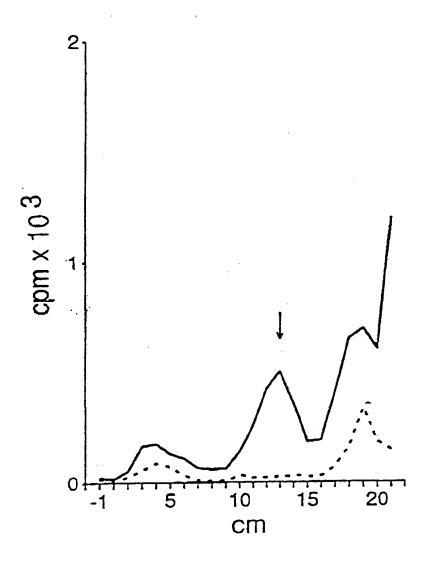


Fig. 8

# INTERNATIONAL SEARCH REPORT

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all)6				
According Int. Cl. <sup>6</sup>	to International Patent classification (IPC) or to both Nation C12N 15/85, 15/60, 15/67	al Classification and IPC		
II. Fi	ELDS SEARCHED			
	Minimum Docum	mentation Searched 7	<del></del>	
Classificat	ion System	Classification Symbols	· · · · · · · · · · · · · · · · · · ·	
IPC W Ch	/PAT Derwent Database: Keywords: inducible, page inducible, pa	promoter, regulatory, element on-coding	t, exon, non-coding	
	Documentation Searched other to the Extent that such Documents are	hen Minimum Documentation e included in the Fields Searched <sup>8</sup>		
Bioteci AU:IPC	hnology Abstracts: Keywords: growth, hormone C:C12N 15/85, 15/60, 15/67, 15/11, 15/18:	e, exon, non-coding		
III. DO	OCUMENTS CONSIDERED TO BE RELEVANT *			
Category*	Citation of Document, <sup>11</sup> with indication, where appropri	iate of the relevant passages 12	Relevant to Claim No 13	
Y	Hampson, R.K. et al. Molecular and Cellular E April 1989 (American Society for Microbiolog "Alternative Processing of Bovine Growth Ho by Downstream Exon Sequences", see pages	gy) rmone mRNA is Influenced	1-7	
Y	Byrne, C.R. et al. Australian Journal of Biolog No. 4, 1987, "The Isolation and Characterisa Hormone Gene", see pages 459-468.	ical Sciences, Volume 40, tion of the Ovine Growth	1-7	
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"A" Document of the control of the c	cial categories of cited documents: 10  ument defining the general state of the art which is considered to be of particular relevance or document but published on or after the national filing date iment which may throw doubts on priority claim(s) hich is cited to establish the publication date of her citation or other special reason (as specified) iment referring to an oral disclosure, use, bition or other means iment published prior t the international filing date after than the priority date claimed	filing date or priority of with the application be principle or theory und document of particula invention cannot be considered to involve "Y" document of particula invention cannot be or invention cannot be or inventive step when the with one or more other combination being obtated in the art.	r relevance; the claimed onsidered novel or cannot be an inventive step relevance; the claimed onsidered to involve an ne document is combined in such documents, such rious to a person skilled in	
IV. CER	TIFICATION	"&" document member of	the same patent family	
<del> </del>	Actual Completion of the International Search	Date of Mailing of this Internation 25 June 1992	onal Search Report	
nternational	ternational Searching Authority Signature of Authorized Officer			
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FU	RTHI	R INFORMATION CONTINUED FROM THE SECOND SHEET
,	A .	Curatola, A.M. and C. Basilico.  Molecular and Cellular Biology, Volume 10, N . 6, June 1980 (American Society for Microbiology)  "Expression of the K-faf Proto-Oncogene Is Controlled by 3 Regulatory Elements Which Are Specific for Embryonal Carcinoma Cells" see pages 2575-2483.  Gutkind, J.S. et al. Molecular and Cellular Biology, Volume 11, No. 3, March 1991 (American Society for Microbiology)  "A Novel c-far Exon Utilized in Epstein-Barr Virus-Infected B Lymphocytes but Not in Normal Monocytes" see pages 1500-1507.
٧.		OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE 1
	interr	ational search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:  Claim numbers, because they relate to subject matter not required to be searched by this Authority, namely:
1.	П	Claim Immuels, Declare the State to State the State that the S
2.		Claim numbers, because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3.		Claim numbers, because they are dependent claims and are not drafted in accordance with the second and third sentences of PCT Rule 6.4s
VI.		OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING 2
This	intern	ational Searching Authority found multiple inventions in this international application as follows:
1. 2.		As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.  As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:
3.		No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:
4.		As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.
Rema		Protest additional search fees were accompanied by applicant's protest.
		rotest accompanied the payment of additional search fees.